#### INVENTOR SEARCH

=> d ibib abs 114 1-1

L14 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1999:576753 HCAPLUS Full-text

DOCUMENT NUMBER: 131:219169

TITLE: Cosmetic or pharmaceutical formulations

containing isoquercetin with antiviral activity
INVENTOR(S): Buchbolz, Herwig; Kraus, Christine; Wagner,

Annette; Meduski, Jerzy

PATENT ASSIGNEE(S): Merck Patent G.m.b.H., Germany

SOURCE: PCT Int. Appl., 23 pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent

LANGUAGE: German FAMILY ACC. NUM. COUNT: 1

FAMILY	ACC.	NUM.	COUNT:	1
PATENT	INFO	RMATI	: MC	

PA	TENT :	NO.			KIN	D	DATE		AP	PLIC	CATI	ON 1	10.		D	ATE	
WO	9944	578			A1		1999	0910	WO	199	99-E	P110	) 4		1	9990	220
	W:	CA,	JP,	US													
	RW:	ΑT,	BE,	CH,	CY,	DE,	DK,	ES,	FI, F	R, (	GΒ,	GR,	ΙE,	IT,	LU,	MC,	NL,
		PT,	SE														
DE	1980	9304			A1		1999	0909	DE	199	98-1	9809	304		1	9980	305
CA	2322	450			A1		1999	0910	CA	199	99-2	3224	150		1	9990	220
EP	1059	911			A1		2000	1220	EP	199	99-9	131	76		1	9990	220
EP	1059	911			В1		2005	0420									
	R:	AT,	BE,	CH,	DE,	FR,	GB,	IT,	LI, N	և, չ	SE,	ΙE					
JP	2002	5052	67		T		2002	0219	JP	200	00-5	3418	31		1	9990	220
AT	2934	33			T		2005	0515	AT	199	99-9	131	76		1	9990	220
US	2002	1515	99		A1		2002	1017	US	199	99-3	4971	13		1	9990	708
PRIORIT	Y APP	LN.	INFO	. :					DE	199	98-1	9809	304	1	A 1	9980	305
									WO	199	99-E	P11(	) 4	1	1	9990	220

- AB Solid or liquid formulations contain isoquercetin as a natural flavonoid. The isoquercetin is contained as a light protection filter and/or an antiviral substance. The invention relates to both cosmetic and pharmaceutical formulations. Thus, a lipstick contained isoquercetin 0.1, Cremophor A-25 20.0, Cetiol HE 22.0, glycerin 5.0, preservative q.s., and water to 100% by weight
- REFERENCE COUNT:
- 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

#### RESULTS FROM REGISTRY AND HCAPLUS

=> d que stat 15

4 SEA FILE=REGISTRY ABB=ON (SILVER OXIDE OR TITANIUM DIOXIDE OR

ZINC OXIDE)/CN

288381 SEA FILE=HCAPLUS ABB=ON L1 OR SILVER OXIDE OR TITANIUM L3

DIOXIDE OR ZINC OXIDE

L444662 SEA FILE=HCAPLUS ABB=ON L3 AND (?PIGMENT? OR ?INK? OR

?LACOUER? OR ?PLASTIC?)

1.5 17 SEA FILE=HCAPLUS ABB=ON L4 AND ?HERPES?

=> d ibib abs 15 1-17

ANSWER 1 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2007:941793 HCAPLUS Full-text DOCUMENT NUMBER: 147:308193

TITLE:

Anti-infective formulation comprising solvent vehicle and solidifying agent

INVENTOR(S):

Zhang, Jie; Warner, Kevin S.; Sharma, Sanjay

PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 21pp., Cont.-in-part of U.S.

Ser. No. 146,917. CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 17

PATENT INFORMATION:

PATENT NO.	KIND	DATE	AP	PLICATION NO.		DATE
US 2007196325	A1	20070823	US	2006-640101		20061214
US 2005276842	A1	20051215	US	2005-146917		20050606
PRIORITY APPLN. INFO.:			US	2004-577536P	P	20040607
			US	2005-146917	A2	20050606
			US	2005-750465P	P	20051214
			US	2005-750522P	P	20051214
			US	2005-750637P	P	20051214
A 20						

The present invention is drawn to solidifying adhesive formulations, methods AB of drug delivery, and solidified layers for dermal delivery of a drug which can treat various skin infections, such as fungal, bacterial, and/or viral skin infections. The formulation can include an anti-infective drug, a solvent vehicle, and a solidifying agent. The solvent vehicle can include a volatile solvent system including at least one volatile solvent, and a nonvolatile solvent system including at least one non-volatile solvent. The nonvolatile solvent system can facilitate the delivery of the drug at therapeutically effective rates for sustained period of time. The nonvolatile solvent system can also act as a plasticizer for the solidifying agent. The formulation can have a viscosity suitable for application to a skin surface prior to evaporation of the volatile solvents system. When applied to the skin, the formulation can form a solidified layer after at least a portion of the volatile solvent system is evaporated Thus, adhesive solidifying formulation was prepared containing ethanol 26%. Eudragit RL PO 44%, isostearic acid 26%, diisopropanol amine 2% and acyclovir 2%.

L5 ANSWER 2 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2007:670138 HCAPLUS Full-text DOCUMENT NUMBER: 147:102133

2

TITLE: Compositions and methods for treating dermatological

conditions

INVENTOR(S): Zhang, Jie; Warner, Kevin S.; Sharma, Sanjay

PATENT ASSIGNEE(S): Zars, Inc., USA SOURCE: PCT Int. Appl., 74pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 17
PATENT INFORMATION:

PA:	TENT :	ΝΟ.			KIN	D	DATE			APPL	ICAT	ION:	NO.				
MO.	2007	0706	43		A 2	_	2007	0621		WO 2	006-	11847	747			0061	
110																	
	W:	ΑĽ,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,
		CN,	co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
		GE,	GH,	GM,	GT,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KM,	KN,
		KP,	KR,	KZ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	LY,	MA,	MD,	MG,	MK,
		MN,	MW,	MX,	MY,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,
		RS,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SM,	SV,	SY,	TJ,	TM,	TN,	TR,	TT,
		TZ,	UA,	UG,	US,	UZ,	VC,	VN,	ZA,	ZM,	ZW						
	RW:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,
		IS,	IT,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,
		CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG,	BW,	GH,
		GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,	BY,
		KG,	ΚZ,	MD,	RU,	TJ,	TM										
PRIORIT:	Y APP	LN.	INFO	. :						US 2	005-	7504	65P		P 2	0051	214
										US 2	005-	7505	22P		P 2	0051	214
										IIS 2	005-	7505	24P		P 2	0051	214
										US 2	005-	1506	3 / 5		P 2	0051	Z14

AB The present invention is drawn to solidifying adhesive formulations, methods of drug delivery, and solidified layers for dermal delivery of a drug which can treat various dermatol. conditions, such as a bacterial infection, a virus infection, a fungal infection, alopecia, dermatitis, psoriasis, or photodamaged skin. The formulation can include a drug, a solvent vehicle, and a solidifying agent. The solvent vehicle can include a volatile solvent system including at least one volatile solvent, and a non-volatile solvent system including at least one non-volatile solvent. The non-volatile solvent system can facilitate the delivery of the drug at therapeutically effective rates for sustained period of time. The non-volatile solvent system can also act as a plasticizer for the solidifying agent. The formulation can have a viscosity suitable for application to a skin surface prior to evaporation of the volatile solvents system. When applied to the skin, the formulation can form a solidified layer after at least a portion of the volatile solvent system is evaporated Thus, an adhesive solidifying formulation was prepared containing acyclovir 3%, ethanol 21%, Eudragit RL-PO 15%, isostearic acid 31%, and trolamine 30%. The formulation provided significant penetration of the active ingredient through hairless mouse and human skin, which was greater than the marketed Zovirax cream. The combination of isostearic acid and trolamine enhanced the flux of acvclovir. The formulation showed a sustained delivery of acyclovir for up to 8 h. It is reasonable to assume based on the drug load and the continued presence of the non-volatile solvent that the delivery of acyclovir would continue at the reported flux values for as long as the subject desires to leave the adhesive solidifying formulation affixed to the skin.

L5 ANSWER 3 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:1256641 HCAPLUS Full-text DOCUMENT NUMBER: 146:50262

TITLE: Antibiotic kit and compositions

INVENTOR(S): Friedman, Doron; Besonov, Alex; Tamarkin, Dov; Eini, Meir

PATENT ASSIGNEE(S): Foamix Ltd., Israel

SOURCE: U.S. Pat. Appl. Publ., 31pp., Cont.-in-part of U.S.

Ser. No. 532,618. CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 21 PATENT INFORMATION:

PATENT 1				NI:	21												
	ENT:				KIN		DATE				ICAT					ATE	
US WO	2006 2004 2004	2694: 0372:	85 25		A1 A2 A3		2006 2004 2004	1130 0506		US 2	006-	4484	90		2	0060	607
	W:	CO, GM, LS, PL,	CR, HR, LT, PT,	CU, HU, LU, RO,	CZ, ID, LV, RU,	DE, IL, MA, SC,	DK, IN, MD,	DM, IS, MG, SE,	DZ, JP, MK, SG,	EC, KE, MN, SK,	BG, EE, KG, MW, SL,	ES, KP, MX,	FI, KR, MZ,	GB, KZ, NO,	GD, LC, NZ,	GE, LK, OM,	GH, LR, PH,
	RW:	GH, KG, FI,	GM, KZ, FR,	KE, MD, GB,	LS, RU, GR,	MW, TJ, HU,	MZ, TM, IE,	SD, AT, IT,	SL, BE, LU,	SZ, BG, MC,	TZ, CH, NL, GW,	CY, PT,	CZ, RO,	DE, SE,	DK, SI,	EE, SK,	ES, TR,
US	2005	0695	66		A1		2005				004-					0040	
	2006				A1		2006				005-					0051	
WO	2007				A2		2007				006-			DV		0060	
	W:										BG,						
											EC, JP,						
											MA,						
											PL,						
											TR,						
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	BM.						CZ.	DE	DK	EE	ES,	FT	FR	GB	GR	нп	TE
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US	2007				A1		2007	1220		US 2	007-	7325	47		2	0070	404
PRIORITY				. :						US 2	002-	4295	46P	1		0021	
										US 2	003-	4923	85P	1	P 2	0030	804
											003-			1		0031	
										US 2	004-	9113	67		A2 2	0040	804
										US 2	005-	6882	44P	1	P 2	0050	607
										US 2	005-	5326	18		A2 2	0051	222
										IL 2	002-	1524	86		A 2	0021	025
										US 2	003-	4976	48P	1	P 2	0030	825
										US 2	003-	5300	15P	1	P 2	0031	216
										US 2	004-	8355	05		A2 2	0040	428
										US 2	004-	9223	58	1	A2 2	0040	820
										US 2	005-	4192	1	1	A2 2	0050	124
										US 2	006-	7891	86P	1	P 2	0060	404
										US 2	006-	4484				0060	607
											006-			1		0061	
										US 2	007-	8804	34P	1	P 2	0070	112

AB The present invention relates to a therapeutic kit to provide an effective dosage of an antibiotic including an aerosol packaging assembly includes a container accommodating a pressurized product; and an outlet capable of releasing the pressurized product as a foam, wherein the pressurized product composition of an antibiotic; at least one organic carrier selected from the group consisting of a hydrophobic organic carrier, an organic polar solvent, an emollient and mixts. at 2-50%, a surfactant, 0.01-5% by weight of at least one polymeric additive selected from the group consisting of a bloadhesive agent, a gelling agent, a film forming agent and a phase change agent, water; and liquefied or compressed gas propellant at 3-25% by weight of the total composition

L5 ANSWER 4 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:1099617 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 145:432172

TITLE: Method for the treatment of herpes zoster

with post-herpetic neuralgia in elderly patients with

organic personality disorder

INVENTOR(S): Arsenenko, L. D.; Arsenenko, A. S.; Sereda, T. V.
PATENT ASSIGNEE(S): Gos. Obrazovatel'noe Uchrezhd. Dopolnitel'nogo Prof.

Obrazovan. Minister. Obrazovan. Ross. Fed.

Novokuznetskii Gos. Inst. Usovershenstvovaniva

Vrachei, Russia Russ., 12pp.

SOURCE: Russ., 12pp.
CODEN: RUXXE7
DOCUMENT TYPE: Patent

LANGUAGE: Russian

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
RU 2285528	C2	20061020	RU 2004-138166	20041227
PRIORITY APPLN. INFO.:			RU 2004-138166	20041227
AB Method is disclosed	for	the treatment	of herpes zoster with	post-herpe

Method is disclosed for the treatment of herpes zoster with post-herpetic neuralgia in elderly patients with organic personality disorder. The invention relates to a method for the treatment of post-herpetic neuralgia in elderly patients with organic disorders of personality. Method involves combination therapy with antiviral agents, immunomodulators, analgesics, nootropic agents, metabolites, cerebroprotective agents, antidepressants and regimen of their administration. Method provides the effective treatment based on the complex effect on different pathogenetic links of the disease and taking into account the specific aging and nervous-psychic features of patients of this group. Method ensures the enhanced effectiveness of treatment.

L5 ANSWER 5 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:1094143 HCAPLUS Full-text

DOCUMENT NUMBER: 145:426012

TITLE: Foamable oil in water emulsion composition comprising

TITLE: Foamable

INVENTOR(S): Tamarkin, Dov; Friedman, Doron; Besonov, Alex; Eini,

Meir

PATENT ASSIGNEE(S): Foamix Ltd., Israel

SOURCE: U.S. Pat. Appl. Publ., 14 pp., Cont.-in-part of U.S. Ser. No. 532,618.

CODEN: USXXCO

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 21

PATENT INFORMATION:

	TENT				KIN		DATE				ICAT					ATE	
	2006				A1		2006				006-					0060	
WO	2004	0372	25		A2		2004			WO 2	003-	IB55	27		2	0031	024
	2004				A3		2004										
	W:	AE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,
							DK,										
							IN,										
							MD,										
							SD,										
							VN,										
	RW:						MZ,					UG.	ZM.	ZW.	AM.	AZ.	BY.
							TM,										
							IE,										
		BF,	BJ,	CF.	CG,	CI,	CM,	GA,	GN,	GO,	GW,	ML,	MR.	NE.	SN,	TD,	TG
US	2005				A1		2005				004-					0040	
ZA	2005	0032			A		2006	0830		ZA 2	005-	3298			2	0050	425
US	2006	1409	84		A1		2006	0629		US 2	005-	5326	18		2	0051	222
AU	2006	2018	78		A1		2007	0927		AU 2	006-	2018	78		2	0060	504
WO	2007	1020	52		A2		2007	0913		WO 2	006-	IB41	70		2	0060	914
WO	2007	1020	52		A3		2008	0103									
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
		GE,	GH,	GM,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KM,	KN,	KP,
		KR,	KZ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	LY,	MA,	MD,	MG,	MK,	MN,
		MW,	MX,	MY,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RS,
		RU,	SC,	SD,	SE,	SG,	SK,	SL,	SM,	SV,	SY,	ΤJ,	TM,	TN,	TR,	TT,	TZ,
		UA,	UG,	US,	UZ,	VC,	VN,	ZA,	ZM,	ZW							
	RW:	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FΙ,	FR,	GB,	GR,	HU,	ΙE,
		IS,	ΙT,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ΒJ,
							GN,										
							NA,					UG,	ZM,	ZW,	AM,	ΑZ,	BY,
		KG,	ΚZ,	MD,	RU,	ΤJ,	TM,	ΑP,	EA,	EP,	OA						
PRIORIT	Y APP	LN.	INFO	.:							002-					0021	
											002-					0021	
											003-					0030	
											003-					0031	
											004-				A2 2		
										US 2	005-	7170	58P			0050	
										US 2	005-	5326	18		A2 2		
										US 2	006-	3897	42		A 2	0060	327

AB The present invention provides a foamable composition for administration to the skin, body surface, body cavity or mucosal surface, e.g., the mucosa of the nose, mouth, eye, ear, respiratory system, vagina or rectum. The foamable oil in water emulsion composition includes: an oil globule system, selected from the group consisting of oil bodies; and sub-micron oil globules, about 0.1% to about 5% by weight of an agent, selected from the group consisting of a surface-active agent, having an HLB value between 9 and 16; and polymeric agent, and a liquefied or compressed gas propellant at a concentration of about 3% to about 25% by weight of the total composition, water and optional ingredients are added to complete the total mass to 100%. Upon release from an aerosol container, the foamable composition forms and expanded foam suitable for topical administration. For example, emulsion composition was prepared comprising mineral oil 5.6%, iso-Pr myristate 5.6%, glyceryl monostearate 0.45%, PEG-40 stearate 2.6%, stearyl alc. 0.85%, Ranthan gum

0.26%, methocel K100M 0.26%, Polysorbate 80 0.90%, water 74.88%, preservative 0.60 and propellant 8%.

L5 ANSWER 6 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:759629 HCAPLUS Full-text

DOCUMENT NUMBER: 145:195180

TITLE: Cosmetic composition comprising hydrophobic and

hydrophilic silica particles

INVENTOR(S): Ingman, Dov

PATENT ASSIGNEE(S): Or-Le-Or Ltd., Israel Eur. Pat. Appl., 30 pp. SOURCE:

CODEN: EPXXDW

Patent DOCUMENT TYPE:

LANGUAGE: English FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	TENT	NO.			KIN	D	DATE			APPL	ICAT	ION I	NO.		D.	ATE		
						-												
EP	1685	824			A1		2006	0802	1	EP 2	006-	1004	58		2	0060	117	
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙT,	LI,	LU,	NL,	SE,	MC,	PT,	
		ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,	PL,	SK,	
		BA,	HR,	IS,	YU													
CA	2534	1306			A1		2006	0730		CA 2	006-	2534	306		2	0060	130	
US	2006	2574	37		A1		2006	1116	1	JS 2	006-	3418	72		2	0060	130	
RTT	Y APP	T.N.	TNFO							TT. 2	005-	1666	46	7	1 2	0050	130	

PRIORITY APPLN. INFO.: IL 2005-166646 The present invention relates to a new topical cosmetic composition formulated for concealing wrinkles and for eliminating or reducing damages to the skin appearance resulted from a wide variety of disorders, such as for example, acne. The composition comprises water, optionally containing 25 to 400 ppm of Ag, hydrophobic particles, preferably hydrophobic silica, having a diameter, ranged from about 5 to about 150 nm, and/or hydrophilic particles, preferably hydrophilic silica, having a diameter, ranged from about 5 to about 150 nm and a soluble electrolyte, capable of releasing free ions in an aqueous environment. Thus, a hypotonic composition for treating acne comprised Dead Sea salt 0.2, zinc sulfate 1, hydrophobic silica 5, hydrophilic silica 5, tea tree oil 2, sea buckthorn oil 3, vitamin A 0.1, vitamin C 1.5, vitamin E acetate 0.1, methylparaben 0.1, propylene glycol 2, and water 80%, resp. A composition comprising water, optionally containing Ag 25 to 400 ppm, 10 weight% Aerosil 380, 2.5 weight% Aerosil R812, 1 to 20 weight% Dead Sea salt, and optionally one or more conventional skincare and/or anti-acne agent, selected from evening primrose oil, sweet almond oil, sea buckthorn oil, tea tree oil, Finsolv TN, (C12-15 alkyl benzoate), octyl hydroxystearate, salicylic acid, vitamin C, citric acid, azelaic acid, benzoyl peroxide, zinc acetate and sulfur. The composition was highly effective in treating acne. The concentration of the salt in such composition was determined according to the treated skin type (dried, oily, etc.) and the particular acne type, grade and state of the treated individual. Compns. containing higher concns. of salt (10 to 20 weight%) are preferred for treating an oily skin and an intensive acne state.

REFERENCE COUNT: THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 7 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:1291998 HCAPLUS Full-text

DOCUMENT NUMBER: 144:40803

TITLE: Vasoactive kit and compositions comprising emollients and polymeric additive

Friedman, Doron; Besonov, Alex; Tamarkin, Dov; Eini, INVENTOR(S):

Meir

PATENT ASSIGNEE(S): Foamix Ltd., Israel

SOURCE: U.S. Pat. Appl. Publ., 18 pp., Cont.-in-part of U.S. Ser. No. 911,367.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 21 PATENT INFORMATION:

		ENT				KIN		DATE				ICAT					ATE	
		2005				A1	_	2005				2005-					0050	
		2004				A2		2004				003-					0031	
		2004				A3		2004								_		
		W:			AT.		AT.			BA.	BB.	BG,	BR.	BY.	B7.	CA.	CH.	CI
												EE,						
												KG,						
												MW,						
												SL,	10,	IM,	IN,	IK,	11,	12
								VN,										
		RW:										TZ,						
												CH,						
												NL,						
			BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	T
1	US	2005	0695	66		A1		2005	0331		US 2	004-	9113	67		2	0040	80.
	ZA	2005	0032	98		A		2006	0830		ZA 2	2005-	3298			2	0050	42
- 2	ΑU	2006	2018	78		A1		2007	0927		AU 2	006-	2018	78		2	0060	50
- 3	ΑU	2006	2832	25		A1		2007	0301		AU 2	006-	2832	25		2	0060	50
1	WO	2007	0233	96		A2		2007	0301		WO 2	006-	IB35	25		2	0060	508
		W:			AL.							BG,			BY.			
												EC,						
												JP,						
												MA,						
												PL,						
				SK,				10,	111,	IN,	ıĸ,	TT,	14,	UM,	UG,	05,	04,	V
				YU,			ZW											
		RW:										ES,						
												RO,						
												MR,						
									SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	B:
			KG,	ΚZ,		RU,												
1	US	2007	0202			A1		2007	0125		US 2	2006-	4889	89		2	0060	719
1	US	2007	2923	59		A1		2007	1220		US 2	2007-	8111	40		2	0070	60'
DR.	IΤ	APP	LN.	INFO	. :						IL 2	2002-	1524	86		A 2	0021	025
											US 2	2002-	4295	46P		P 2	0021	129
											US 2	003-	4923	85P		P 2	0030	80.
											WO 2	003-	IB55	27		W 2	0031	02
											US 2	2004-	9113	67		A2 2	0040	80.
												2003-					0030	
												003-					0031	
												2004-					0040	
												2004-					0040	
												2005-					0050	
												2005-					0050	
												2005-					0050	
												005-					0050	
												2005-					0051	
											HS 2	006- 006-	7818	68P		P 2	0060	31
														002			0060	

US 2006-811627P P 20060607 US 2006-481596 A2 20060706 US 2006-488989 A2 20060719 US 2007-897638P P 20070126 US 2007-899176P P 20070202 US 2007-717897 A2 20070313

AB The present invention relates to a therapeutic kit to provide an effective dosage of a vasoactive agent, including an aerosol packaging assembly with a container accommodating a pressurized product; and an outlet capable of releasing the pressurized product as a foam. The pressurized product comprises a foamable composition including: a vasoactive agent; a carrier selected from the group consisting of a hydrophobic organic carrier, an organic polar solvent, an emollient and mixts: thereof at 2-50% by weight, a surfactant, 0.01-5% by weight of at least 1 polymeric additive selected from the group consisting of a bioadhesive agent, a gelling agent, a film forming agent and a phase change agent, water; and liquefied or compressed gas propellant at a concentration of 3-25% by weight of the total composition

L5 ANSWER 8 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:904090 HCAPLUS Full-text

DOCUMENT NUMBER: 143:235474

TITLE: Cosmetic and pharmaceutical foam with solid particles

such as oxides for topical administration

INVENTOR(S): Tamarkin, Dov; Friedman, Doron; Eini, Meir; Besonov,

Alex

PATENT ASSIGNEE(S): Foamix Ltd., Israel

SOURCE: U.S. Pat. Appl. Publ., 18 pp.

CODEN: USXXCO DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	US 2005186147	A1	20050825	US 2005-50999	20050204
	AU 2005201455	A1	20050825	AU 2005-201455	20050204
PRI	ORITY APPLN. INFO.:			US 2004-541698P P	20040204
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AB The invention relates to an alc.-free cosmetic or pharmaceutical foam carrier comprising about 2 to 30% by weight solid particles, about 2 to 75% by weight hydrophobic solvent, about 10 to 85% by weight water, about 0.1 to 5% by weight surface-active agent, about 0.1 to 5% by weight stabilizer/gelling agent and a liquefied or compressed gas propellant in a container, which upon release provides a breakable foam suitable for topical administration. For example, a wound healing foam was prepared containing mineral oil 12.5, colloidal silver 2.0, lidocaine 4.0, Arlacel 135 2.0, Avicel CL611 2.0, Tween 80 2.0, cocoamidopropylbetaine 1.0, D-Panthenol 50P 10.0, benzalkonium chloride 0.20 and water to 100%.

L5 ANSWER 9 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:904087 HCAPLUS Full-text

DOCUMENT NUMBER: 143:235471

TITLE: Kit and composition of imidazole with enhanced bioavailability and therapeuticc uses thereof

INVENTOR(S): Tamarkin, Dov; Friedman, Doron; Eini, Meir

PATENT ASSIGNEE(S): Foamix Ltd., Israel

SOURCE: U.S. Pat. Appl. Publ., 19 pp., Cont.-in-part of U.S.

Ser. No. 911,367.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 21

PATENT INFORMATION:

PATENT NO. DATE KIND DATE APPLICATION NO. --- -----A1 20050825 US 2005-41921 US 2005186142 WO 2004037225 A2 20040506 WO 2003-IB5527 WO 2004037225 A3 20041229 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG US 2005069566 A1 20050331 US 2004-911367 20040804 ZA 2005003298 A 20060830 ZA 2005-3298 20050425 2A 2005003298 A 20060630 2A 200753290 20050830 2A 200575290 2A 20057529 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN. YU, ZA, ZM, ZW RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM AU 2006201878 A1 20070927 US 2007292355 A1 20071220 IN 2007CN03681 A 20071116 AU 2006-201878 20060504 US 2007-732547 20070404 IN 2007-CN3681 20070823 IL 2002-152486 A 20021025 US 2003-492385P P 20030804 AU 2006-201878 20060504 PRIORITY APPLN. INFO.: WO 2003-TB5527 A 20031024 US 2004-911367 A2 20040804 US 2002-429546P P 20021129 P 20030825 US 2003-497648P P 20031216 US 2003-530015P US 2004-835505 A2 20040428 A2 20040820 US 2004-922358 US 2005-41921 A 20050124 US 2005-688244P P 20050607 US 2005-688244P P 20050607 US 2005-532618 A2 20051222 US 2006-789186P P 20060124 US 2006-789186P P 20060044 US 2006-448490 A2 20060607 US 2006-861620P P 20061129 US 2007-880434P P 20070112

AB The present inventino relates to a composition and therapeutic kit comprising therapeutic azole with increased solubility. The kit includes an aerosol

packaging assembly containing a container accommodating a pressurized product and an outlet capable of releasing the pressurized product as a foam. The pressurized product includes a foamable composition including: i. a therapeutic azole, wherein the solubility of the azole in the composition before foaming is less than the solubility of the azole in the composition after foaming; ii. at least one organic carrier selected from the group consisting of a hydrophobic organic carrier, a co-solvent, an emollient and mixts. thereof, at a concentration of about 2% to about 5% by weight; iii. a surface-active agent; iv. about 0.01% to about 5% by weight of at least one polymeric additive selected from the group consisting of a bloadhesive agent, a gelling agent, a film forming agent and a phase change agent; v. water; and vi. liquefied or compressed gas propellant at a concentration of about 3% to about 25% by weight of the total composition

L5 ANSWER 10 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:902663 HCAPLUS Full-text

DOCUMENT NUMBER: 143:235459

TITLE: Cosmetic and pharmaceutical foam with solid particles

such as oxides for topical administration

INVENTOR(S): Tamarkin, Dov; Friedman, Doron; Eini, Meir; Besonov,

Alex

PATENT ASSIGNEE(S): Foamix Ltd., Israel SOURCE: PCT Int. Appl., 46 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

	TENT				KIN						ICAT					ATE		
WO	2005 2005	0766	97		A2		2005	0825										
	W:							AZ,										
								DK,										
								IL,										
								MA,										
		NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
		ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW,	SM
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	
		AZ,	BY,	KG,	KZ,	MD,	RU,	ТJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	
		EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,	IS,	IT,	LT,	LU,	MC,	NL,	PL,	PT,	
		RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	
		MR,	NE,	SN,	TD,	TG,	AP,	EA,	EP,	OA								
AU	2005	2014	55		A1		2005	0825		AU 2	005-	2014	55		2	0050	204	
	2555															0050		
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TN	2006						2007	0601		TNI 2	006-	MATO S	12		2	0060	0.01	
PRIORIT							2007	0001			004-							
PRIORII	1 APP	LIN.	INFO	. :														
										WU Z	005-	1012	21		w Z	0050	204	

AB The invention relates to an alc.-free cosmetic or pharmaceutical foam carrier comprising about 2 to 30% by weight solid particles, about 2 to 75% by weight hydrophobic solvent, about 10 to 85% by weight water, about 0.1 to 5% by weight surface-active agent, about 0.1 to 5% by weight stabilizer/gelling agent and a liquefied or compressed gas propellant in a container, which upon release provides a breakable foam suitable for topical administration. For

example, a wound healing foam was prepared containing mineral oil 12.5, colloidal silver 2.0, lidocaine 4.0, Arlacel 135 2.0, Avicel CL611 2.0, Tween 80 2.0, cocoamidopropylbetaine 1.0, D-Panthenol 50P 10.0, benzalkonium chloride 0.20 and water to 100%.

ANSWER 11 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:878561 HCAPLUS Full-text

DOCUMENT NUMBER: 141:346084

TITLE: Device for analysis or separation containing an active

nanostructured carrier, its preparation method and

applications INVENTOR(S): Zou, Fanglin; Chen, Chunsheng; Chen, Ning; Wang,

Jianxia

PATENT ASSIGNEE(S): Chengdu Kuachang Medical Industrial Limited, Peop. Rep. China; Chengdu Kuachang Science & Technology Co.,

Ltd

PCT Int. Appl., 62 pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE: Pat.ent. LANGUAGE: Chinese

FAMILY ACC. NUM. COUNT: 8 PATENT INFORMATION:

PA:	TENT I	NO.			KIN	D	DATE			APPL					D	ATE		
WO	2004	0905	48		A1		2004			WO 2					2	0040	315	
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,	
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,	
		NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
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		BY,	KG,	KΖ,	MD,	RU,	ΤJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	
		ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	
		SK,	TR,	BF,	BJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	
		TD,	TG															
CN	1434	295			A		2003	0806		CN 2					2	0030.	313	
CN	1514	243			A		2004	0721		CN 2	003-	1177	87		2	0030	430	
ΝO	2004	0815	71		A1		2004	0923		WO 2	004-0	CN77			2	0040	120	
	W:	ΑE,	AG,	ΑL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,	
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	ΚP,	KR,	ΚZ,	LC,	
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NA,	ΝI,	
		NO,	ΝZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
		TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	zw	
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		BY,	KG,	ΚZ,	MD,	RU,	ΤJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	
		ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	
		TR,	BF,	ΒJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	1
WO	2004				A1		2004			wo 2						0040		
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							DE,											
							ID,											
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NA,	NI,	
		NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
		TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	zw	
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	
		AZ,	BY,	KG,	KZ,	MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	

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EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
           SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
           SN. TD. TG
    CN 1697975
                       Α
                             20051116 CN 2004-80000653
                                                            20040430
    EP 1624306
                             20060208
                                       EP 2004-730459
                                                             20040430
                       A1
        R: DE, FR, GB, IT
    JP 2007502998
                       T
                           20070215 JP 2006-529552
                                                             20040430
    US 2006057631
                      A1 20060316
                                       US 2005-258996
                                                             20051027
                                                        A 20030313
A 20030430
PRIORITY APPLN. INFO.:
                                        CN 2003-117446
                                        CN 2003-117787
                                        WO 2004-CN77
                                                         A 20040120
                                        WO 2004-CN203
                                                          A 20040315
                                        WO 2004-CN437
                                                         W 20040430
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AB The invention relates to an active nanostructured carrier with high sensitivity for separation and/or anal., its preparation method, and a nanolabel and its labeling method. The invention also relates to a biochip and a polypeptide detection device, all of which contain the high-sensitive active nanostructured carrier and /or the nano-label, especially anal. chips, enzyme labeling plates and chromatoc. test strips, and their applications.

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 12 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2004:352956 HCAPLUS Full-text

DOCUMENT NUMBER: 140:363037

TITLE: Formulations for topical delivery of bioactive

substances and methods for their use

INVENTOR(S): Vromen, Jacob

PATENT ASSIGNEE(S): Australian Importers Ltd., USA SOURCE: U.S. Pat. Appl. Publ., 11 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PRI

PATENT NO.					KIN	D	DATE			APPLICATION NO.				DATE				
	US 2004081681 US 7241456													20021025				
							2007											
CA	2543	370			A1		20040513			CA 2	003-	2543	370	20031015				
WO	2004	0393	48		A1		2004	0513		WO 2	003-	US32	638		2	0031	015	
	W:	AE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,	
		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	
		GM.	HR.	HU.	ID.	IL.	IN,	IS.	JP.	KE.	KG.	KP.	KR.	KZ.	LC.	LK.	LR.	
							MD,											
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	RW:						MZ,							7.W.	AM.	AZ.	BY.	
							TM.											
							IE,											
							CM,											
														20031015				
EP	1558																	
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,	
		IE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,	SK		
US	2007	0717	11		A1		2007	0329		US 2	006-	5352	13		2	0060	926	
	Y APP										002-							
											003-							
										NO Z	005-	0002	050		n 2	OOST	013	

AB The invention relates to topical delivery of bioactive agents. More particularly, the invention relates to anhydrous formulations for percutaneous absorption. The invention provides formulations that allow efficient topical delivery of high concns. of bioactive substances for percutaneous absorption. The formulations according to the invention are generally non-irritating to the skin. A preferred topical formulation comprises (1) anhydrous media containing glycerin, propylene glycol, capric/caprylic triglyceride, cetearyl alc., d-tocopherol, ascorbyl palmitate, thiodipropionic acid, BHT, phenoxyethanol, and parabens and (2) bioactive substances containing micronized niacinamide, micronized acetylsalicylic acid, and micronized ascorbic acid.

REFERENCE COUNT:

45 THERE ARE 45 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 13 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2002:595341 HCAPLUS Full-text

DOCUMENT NUMBER: 137:159019

TITLE: Products for topical applications comprising oil

bodies

INVENTOR(S): Deckers, Harm M.; Van Rooijen, Gijs; Boothe, Joseph;

Goll, Janis; Moloney, Maurice M.

PATENT ASSIGNEE(S): Sembiosys Genetics Inc., Can.

SOURCE: U.S. Pat. Appl. Publ., 27 pp., Cont.-in-part of U.S. Ser. No. 577,147.

CODEN: USXXCO

DOCUMENT TYPE: Patent
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 8

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE		
US 2002106337	A1	20020808	US 2001-983546		20011024		
US 6599513	B2	20030729					
IN 190035	A1	20030531	IN 1998-DE1401		19980525		
ZA 9804459	A	19990413	ZA 1998-4459		19980526		
US 6146645	A	20001114	US 1998-84777		19980527		
US 6183762	B1	20010206	US 1999-448600		19991124		
CA 2290278	A1	20010524	CA 1999-2290278		19991124		
CA 2290278	С	20030729					
US 6372234	B1	20020416	US 2000-577147		20000524		
AU 772919	B2	20040513	AU 2001-85511		20011029		
IN 195229	A1	20050128	IN 2002-DE1185		20021125		
PRIORITY APPLN. INFO.:			US 1997-47753P	P	19970527		
			US 1997-47779P	P	19970527		
			US 1998-75863P	P	19980225		
			US 1998-75864P	P	19980225		
			US 1998-84777	A2	19980527		
			US 1999-448600	A2	19991124		
			US 2000-577147	A2	20000524		
			IN 1998-DE1401		19980525		
			AU 1998-75178		19980527		
			HO 1550 /51/0	110	13300327		

AB The present invention provides novel emulsion formulations which comprise oil bodies. The invention also provides a method for preparing the emulsions and the use of the emulsions in products for topical application to the skin. The products are very mild to the skin and may be easily formulated into a wide variety of personal care and dermatol. products. A stabilized oil body formulation contained Safflower oils 96.50, Keltrol CG 0.70, Arlacel-165 2.50, phytic acid 0.10, and Glydant Plus 0.20%. A low detergent active body wash formulation comprised cetyl hydroxyethyl cellulose 1.00, Miracare BT 5.00,

lauramide DEA 3.00, glycerin 3.00, Na2EDTA 0.05, Polysorbate-20 0.5, Glydant Plus 0.1, lanolin alc. 1.00, petrolatum 3.00, 30% ammonium lauryl sulfate 15.0, the above stabilized oil bodies 25.0, and citric acid 0.89%, water and fragrance qs.

L5 ANSWER 14 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2002:555377 HCAPLUS Full-text

DOCUMENT NUMBER: 137:99039

TITLE: Stabilized brivadine topical formulations containing

oxide pigments

INVENTOR(S): Gehlert, Ulrike; Groeger, Karsten; Schmitz, Reinhard;

Schrader, Karl-Heinz; Schrader, Andreas; Wihsmann, Marc; Maggi, Carlo Alberto; Manzini, Stefano;

Stubinski, Bettina

PATENT ASSIGNEE(S): Berlin-Chemie A.-G., Germany; Menarini Ricerche S.p.A.

SOURCE: PCT Int. Appl., 21 pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent

DOCUMENT TYPE: LANGUAGE:

LANGUAGE: English FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

										APPLICATION NO.										
						A2														
Ţ.	IO.	2002	0569	13		A3		20021107												
		W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BE	BG,	BR,	BY,	BZ,	CA,	CH,	CN,		
			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC	, EE	ES,	FI,	GB,	GD,	GE,	GH,		
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE	, KG	KP,	KR,	KZ,	LC,	LK,	LR,		
			LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN	I, MW	MX,	MZ,	NO,	NZ,	OM,	PH,		
			PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK	, SL	TJ,	TM,	TN,	TR,	TT,	TZ,		
			UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZM,	ZW	7								
		RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ	, TZ	UG,	ZM,	ZW,	ΑT,	BE,	CH,		
								FR,												
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(	ĊΑ	2434	743			A1		2002	0725		CA	2002	-2434	743		2	0020	110		
P	ΔŲ	2002244642			A1		2002	0730		AU	2002	-2446	42		2	0020	110			
		200300322																		
		2003									HU	2003	-2741			2	0020	110		
								2007												
E	EΡ									EP 2002-712810										
		R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GF	R, IT	LI,	LU,	NL,	SE,	MC,	PT,		
			ΙE,	SI,	LT,	LV,	FI,	RO,												
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		1079				A		2004									0030			
		20031						2003				2003					0030			
		2003				A		2003				2003					0030			
								2004			ZA	2003	-5437				0030			
						A1		2004	0506			2003					0031			
PRIORI	IΤ	APP:	LN.	INFO	.:							2001								
											WO	2002	-EP16	3		W 2	0020	110		

AB The use of metal oxide pigments as photodegrdn. stabilizers in topical compns. containing brivudine. is disclosed. Thus, a hydrogel contained brivudine 5, TiO2 21.25, Al2O3 1.25, silica 0.125, glycerol 2.375, iron oxide yellow 3.25, iron oxide red 1.25, iron oxide black 0.5, propylene glycol 20, Paraffinum

subliquidum 5, iso-Pr myristate 5, cetyl alc. 3, polyoxyethylene monostearate 0.8, hydroxyethyl cellulose 0.3, citric acid gs and water to 100 g.

ADDITION NO.

DATE

L5 ANSWER 15 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1998:293380 HCAPLUS Full-text

EIND DATE

DOCUMENT NUMBER: 128:312936

TITLE: Antiherpetic topical pharmaceutical compositions

containing acyclovir

INVENTOR(S): Santus, Giancarlo; Golzi, Roberto; Garavaglia, Antonio PATENT ASSIGNEE(S): Recordati S.A. Chemical and Pharmaceutical Company,

Italy

SOURCE: PCT Int. Appl., 26 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.											APPLICATION NO.								
										WO 1997-EP6022									
												, BY,							
												, IL,							
			KZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MD	, MG,	MK,	MN,	MW,	MX,	NO,	NZ,	
			PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK	, SL,	TJ,	TM,	TR,	TT,	UA,	UG,	
			UZ,	VN,	YU,	ZW													
		RW:	GH,	KE,	LS,	MW,	SD,	SZ,	UG,	ZW,	AT	, BE,	CH,	DE,	DK,	ES,	FI,	FR,	
			GB,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE	, BF,	BJ,	CF,	CG,	CI,	CM,	GA,	
			GN,	ML,	MR,	NE,	SN,	TD,	TG										
	CA	2270	254			A1		1998	0507		CA	1997-	2270	254		1	9971	031	
	AU	9869	078			A	1998	0522	AU 1998-69078						1	9971	031		
	AU	7247	41			B2		2000	0928										
	EP	9483	32			A1		1999	1013		EP	1997-	9488	49		1	9971	031	
	EP	9483	32			B1		2003	0903										
		R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR	, IT,	LI,	LU,	NL,	SE,	MC,	PT,	
			IE,	SI,	LT,	LV,	FI,	RO											
	CN	1235	547			A		1999	1117		CN	1997-	1993	17		1	9971	031	
	BR	9712	610			A		1999	1221		BR	1997-	1261	0		1	9971	031	
	NZ	3356	85			A		2000	0327		NZ	1997-	3356	85		1	9971	031	
		2001		04		T		2001				1998-					9971		
		2485						2003				1997-					9971		
	KR	2000	0526	78		A		2000	0825			1999-					9990	420	
	NO	9901	910			A		1999	0621			1999-					9990	421	
		9903				A		2000	0228			1999-					9990		
	PRIORIT:	Y APP	LN.	INFO	. :							1996-					9961		
											WO	1997-	EP60	22		W 1	9971	031	

AB Described are antiherpetic pharmaceutical compons. suitable for administration by means of a topical applicator (a stick or a medicated roll-on stick). These compons. contain acyclovir (I) or any of its derivs., either alone or associated with vitamin A or any of its esters, as an active ingredient, and are useful in particular for the treatment of herpes labialis. Thus, 5% micronized I was dispersed under stirring at 60° up to homogeneous dispersion in a fatty phase made up of carnauba wax 10, beeswax 15, lanolin 5, cetyl alc. 5, hydrogenated castor oil 60%, the melted mass containing I was then filled into 3 gal cylindrical containers for sticks. The efficacy and comfort of use of the composition in 30 patients with herpes labialis is reported.

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 16 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1997:195630 HCAPLUS Full-text

DOCUMENT NUMBER: 126:190940

TITLE: Topical pharmaceutical compositions containing volatile oils, silicones, and active ingredients INVENTOR(S): Grollier, Jean-Francois; Allec, Josiane; Agostini,

Isabelle

PATENT ASSIGNEE(S): L'Oreal S. A., Fr. SOURCE: Eur. Pat. Appl., 11 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent. LANGUAGE: French

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 755675	A1	19970129	EP 1996-401491	19960705
EP 755675	B1	19970917		
R: DE, ES, FR,	GB, IT			
FR 2737118	A1	19970131	FR 1995-9252	19950728
FR 2737118	B1	19970905		
ES 2109832	T3	19980116	ES 1996-401491	19960705
AU 9659430	A	19970213	AU 1996-59430	19960710
AU 679663	B2	19970703		
CA 2182226	A1	19970129	CA 1996-2182226	19960726
JP 09040548	A	19970210	JP 1996-198146	19960726
JP 2965510	B2	19991018		
US 6136332	A	20001024	US 1996-688027	19960729
PRIORITY APPLN. INFO.:			FR 1995-9252	19950728
OTHER SOURCE(S):	MARPAT	126:190940		

AR The title pharmaceutical containing volatile oils, Ph silicones, and active ingredients are claimed. A pliable paste contained cyclopentadimethylsiloxane 45, polyphenylmethylsiloxane 25, silicone wax 10, polyethylene wax 5, alkyl dimethicone 5, titanium dioxide 5, Nylon 3, and fusidic acid 2q.

L5 ANSWER 17 OF 17 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1994:239633 HCAPLUS Full-text

DOCUMENT NUMBER: 120:239633

TITLE: Devices and methods for detection of an analyte based

upon light interference

INVENTOR(S): Bogart, Gregory R.; Moddel, Garret R.; Maul, Diana M.; Etter, Jeffrey B.; Crosby, Mark; Miller, John B.;

Blessing, James; Kelley, Howard; Sandstrom, Torbjorn;

Stiblert, Lars

PATENT ASSIGNEE(S): Biostar, Inc., USA SOURCE: PCT Int. Appl., 208 pp. CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 14 PATENT INFORMATION:

PA:	CENT I	NO.			KINI	)	DATE			APP:	LICAT:	I NOI	NO.		D2	ATE	
						-											
WO	9403	774			A1		1994	0217		WO :	1993-l	JS56	73		19	9930	610
	W:	AT,	AU,	CA,	JP												
	RW:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR	, IE,	IT,	LU,	MC,	NL,	PT,	SE
AU	9179	004			A		1992	1021		AU :	1991-	7900	4		19	9910	320

2.77	CE 20 40		no.	19941020				
						4004 040056	19910320	
						1991-910026	19910320	
EP	539383			19960918				
				FR, GB, IT,				
JP	05506936		T	19931007	JP	1991-509344	19910320	
JP	3193373		B2	20010730				
ES	2094224		Т3	19970116	ES	1991-910056	19910320	
JP	2001235473	3	A	20010831	JP	2000-287242	19910320 19910320 19930610 19930610	
AU	9345360		A	19940303	AU	1993-45360	19930610	
JP	07509565		T	19951019	JP	1994-505280	19930610	
				20040315				
						1993-915341	19930610	
EP	727038		B1					
	R: ES, I	FR, GB,	IT,	SE				
	1126278		A2	20010822 20011017	EP	2001-108521	19930610	
	1126278		A3	20011017				
EP	1126278		B1					
	R: ES, I	FR, GB,	IT,	SE				
JP	200211620	8	A	20020419 20040315	JP	2001-236186	19930610	
JP	3507048		B2	20040315				
JP	200213949	В	A	20020517	JP	2001-236144	19930610	
				20031014				
EP	1635162		A2	20060315	EP	2005-27236	19930610	
	R: AT, I	BE, CH,	DE,	DK, ES, FR,	GB, G	R, IT, LI, LU,	NL, SE, MC, PT,	ΙE
HK	1003304		A1	20041013	HK	1998-102353	19980319 20010803	
JP	200212260	1	A	20020426 20050720	JP	2001-236166	20010803	
JP	3673849		B2	20050720				
JP	2002122603	3	A	20020426	JP	2001-236198	20010803	
	3547723		B2	20040728				
JP	2005049356	6	A	20050224	JP	2004-276389	20040924	
				20060621				
	Y APPLN. II				US	1992-924343	A 19920731	
					EP	1991-910056	A 19910320	
					JP	1991-509344	A3 19910320	
					WO	1991-910056 1991-509344 1991-US1781	A 19910320	
					EP	1993-915341	A3 19930610	
							A3 19930610	
							W 19930610	
							A3 20010803	
		_			J.F	2001-230100	A3 20010003	

AB Methods for analyzing an optical surface for an analyte of interest in a test sample and related instruments/devices are disclosed. The method entails the use of a thin-film optical immunoassay device whereby an analyte of interest is detected in a test sample through spectral changes in the light impinging on the surface prior to and after the binding of the analyte to a reactive substrate layer(s). The device includes a substrate which has a 1st color in response to light impinging thereon. The substrate also exhibits a 2nd color which is different from the 1st color. The 2nd color is exhibited in response to the same light when the analyte is present on the surface. Thus, SiO was vapor deposited on a polished monocryst. Si wafer to a thickness of 550 Å; the film had a golden interference color. The film was activated with N-(2aminoethyl)-3-aminopropyltrimethoxysilane, coated with a DNP-albumin conjugate to a thickness of  $40\text{\AA}$ , rinsed, and dried. The coated wafer was used in a competitive immunoassay for DNP using goat anti-DNP antibody and an ellipsometer to measure the change in mass at the surface from the change in light intensity.

RESULTS FROM MEDLINE, BIOSIS, KOSMET, RAPRA, EMBASE, AND DRUGU

=> => d que stat 118

4 SEA FILE=REGISTRY ABB=ON (SILVER OXIDE OR TITANIUM DIOXIDE OR

ZINC OXIDE)/CN

24737 SEA L1 OR SILVER OXIDE OR TITANIUM DIOXIDE OR ZINC OXIDE L16

L17 5274 SEA L16 AND (?PIGMENT? OR INK? OR ?LACQUER? OR ?PLASTIC?)

L18 6 SEA L17 AND ?HERPES?

=> d ibib abs 118 1-6

L18 ANSWER 1 OF 6 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on STN

ACCESSION NUMBER: 2006:202468 BIOSIS Full-text

DOCUMENT NUMBER: PREV200600204867

TITLE: LMP1 signaling and activation of NF-kappa B in LMP1

transgenic mice.

AUTHOR(S): Thornburg, N. J.; Kulwichit, W.; Edwards, R. H.; Shair, K.

H. Y.; Bendt, K. M.; Raab-Traub, N. [Reprint Author] CORPORATE SOURCE: Univ N Carolina, Lineberger Comprehens Canc Ctr, Dept

Immunol Microbiol, CB 7295, Room 102, Mason Farm Rd, Chapel

Hill, NC 27599 USA

nrt@med.unc.edu

Oncogene, (JAN 2006) Vol. 25, No. 2, pp. 288-297. SOURCE:

CODEN: ONCNES. ISSN: 0950-9232.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 22 Mar 2006

Last Updated on STN: 22 Mar 2006

Transgenic mice expressing Epstein-Barr virus (EBV) latent membrane protein 1 AB (LMP1) under the control of an immunoglobulin heavy-chain promoter and

enhancer develop lymphoma at a threefold higher incidence than LMP1-negative mice. In vitro, LMP1 activates numerous signaling pathways including p38, c-Jun N terminal kinase (JNK), phosphatidylinositol 3 kinase (PI3K)/Akt, and NFkappa B through interactions with tumor necrosis receptor-associated factors (TRAFs). These pathways are frequently activated in EBV-associated malignancies, although their activation cannot be definitively linked to LMP1 expression in vivo. In this study, interactions between LMP1 and TRAFs and the activation of PI3K/Akt, JNK, p38, and NF-kappa B were examined in LMP1 transgenic mice. LMP1 co-immunoprecipitated with TRAFs 1, 2, and 3. Akt, JNK, and p38 were activated in LMP1-positive and -negative splenocytes as well as LMP1-positive and -negative lymphomas. Multiple forms of NF-kappa B were activated in healthy splenocytes from LMP1 transgenic mice, in contrast to healthy splenocytes from LMP1-negative mice. However, in both LMP1-positive and -negative lymphomas, only the oncogenic NF-kappa B c-Rel, was specifically activated. Similarly to EBV-associated malignancies, p53 protein was detected at high levels in the transgenic lymphomas, although mutations were not detected in the p53 gene. These data indicate that NF-kappa B is activated in LMP1-positive healthy splenocytes; however, NF-kappa B c-Rel is specifically activated in both the transgenic lymphomas and in the rare lymphomas that develop in negative mice. The LMP1-mediated activation of NF-kappa B may contribute to the specific activation of c-Rel and lead to the increased development of lymphoma in the LMP1 transgenic mice.

L18 ANSWER 2 OF 6 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on STN ACCESSION NUMBER: 2000:124940 BIOSIS Full-text

DOCUMENT NUMBER: PREV200000124940

TITLE: A dermatology ward at the beginning of the 20th century.

Albert, Michael R. [Reprint author]; Mackool, Bonnie T. AUTHOR(S): CORPORATE SOURCE: Dermatology Branch, National Cancer Institute, NIH, 10 Center Dr. Building 10, 12N262, Bethesda, MD, 20892-1908,

USA SOURCE:

Journal of the American Academy of Dermatology, (Jan., 2000) Vol. 42, No. 1 Part 1, pp. 113-123. print.

ISSN: 0190-9622.

DOCUMENT TYPE: Article

General Review; (Literature Review)

LANGUAGE: English

ENTRY DATE: Entered STN: 5 Apr 2000

Last Updated on STN: 3 Jan 2002

L18 ANSWER 3 OF 6 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on STN

ACCESSION NUMBER: 1997:54500 BIOSIS Full-text

DOCUMENT NUMBER: PREV199799353703

TITLE: Increased expression of NF-kappa-B proteins P65, P50 and

c-Rel in immunodeficiency-related EBV positive non

Hodgkin's lymphomas.

AUTHOR(S): Feuillard, J.; Martin, A.; Asso-Bonnet, M.; Davi, F.; El

Mansouri, S.; Raphael, M.

CORPORATE SOURCE: Hematol. Surgical Pathol., Avicenne Hospital, UFR, SMBH Bobigny, URA CNRS 625, Pitie Salpetriere, Paris, France

SOURCE: Blood, (1996) Vol. 88, No. 10 SUPPL. 1 PART 1-2, pp. 380A. Meeting Info.: Thirty-eighth Annual Meeting of the American

Society of Hematology, Orlando, Florida, USA, December

6-10, 1996.

CODEN: BLOOAW. ISSN: 0006-4971.

DOCUMENT TYPE: Conference; (Meeting)

Conference; Abstract; (Meeting Abstract)

Conference; (Meeting Poster)

LANGUAGE: Enalish

ENTRY DATE: Entered STN: 4 Feb 1997 Last Updated on STN: 5 Feb 1997

L18 ANSWER 4 OF 6 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights

reserved on STN

ACCESSION NUMBER: 2006576090 EMBASE Full-text

Incorporating Photodynamic Therapy into a Medical and TITLE:

Cosmetic Dermatology Practice.

AUTHOR: Gilbert D.J.

CORPORATE SOURCE: Dr. D.J. Gilbert, Newport Dermatology and Laser Associates,

1441 Avacado, Newport Beach, CA 92660, United States. lazrdoc@pacbell.net

SOURCE: Dermatologic Clinics, (Jan 2007) Vol. 25, No. 1, pp.

111-118.

Refs: 17

ISSN: 0733-8635 CODEN: DRMCDJ

PUBLISHER IDENT .: S 0733-8635(06)00100-8

United States COUNTRY:

DOCUMENT TYPE: Journal; General Review; (Review)

FILE SEGMENT: 013 Dermatology and Venereology

030 Clinical and Experimental Pharmacology

Health Policy, Economics and Management 036

037 Drug Literature Index

Adverse Reactions Titles 038

LANGUAGE: English

ENTRY DATE: Entered STN: 12 Dec 2006

Last Updated on STN: 12 Dec 2006

AB ALA-PDT is a safe, effective, and easy-to-perform procedure for the treatment of a variety of cutaneous conditions. Pretreatment and posttreatment procedures are straightforward and well documented. Adverse effects are mild, temporary, and easily managed. Light sources may already be available in physicians' offices or may be purchased for \$8000. Cosmetic benefits of ALA-PDT encourage patients to seek additional cosmetic treatments, increasing practice revenue. .COPYRGT. 2006 Elsevier Inc. All rights reserved.

L18 ANSWER 5 OF 6 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights

reserved on STN

ACCESSION NUMBER: 1996129090 EMBASE Full-text

TITLE: [Sun-induced skin damage].

SCHADWIRKUNGEN DER SONNENSTRAHLUNG AN DER HAUT.

AUTHOR: Raab W.

CORPORATE SOURCE: Prof. Dr. W. Raab, Walfischgasse 3, A-1010 Wien 1, Austria SOURCE: Aktuelle Dermatologie, (1996) Vol. 22, No. SUPPL. 1, pp.

2-6.

ISSN: 0340-2541 CODEN: AKDEDY

COUNTRY: Germany

DOCUMENT TYPE: Journal; Conference Article; (Conference paper)

FILE SEGMENT: 013 Dermatology and Venereology
037 Drug Literature Index

LANGUAGE: German

SUMMARY LANGUAGE: English; German

ENTRY DATE: Entered STN: 11 Jun 1996

Last Updated on STN: 11 Jun 1996

AB Ultraviolet-induced skin lesions may be provoked via different mechanisms. Most frequently, an overload of the natural sun defense is encountered, resulting in sunburn - seen immediately - or in chronic sun damage emerging only after decades. With the increasing life expectancy, the symptoms of such a chronic sun damage are more and more often noticed by the dermatologist (skin dryness, premature skin ageing, pigmented spots, actinic keratoses, nonmelanoma skin cancer). Other sun-induced skin lesions include specific dermatoses caused by ultraviolet rays ('sun-Kobner'), the consequences of immunosuppression (Herpes simplex 'solaris', LE, melanoma) and photodynamic reactions due to, for example, various drugs. True photodermatoses are only rarely seen, possibly due to the fact that their diagnosis is rather complicated and mild cases are misdiagnosed. - Sun protection in general and even more in the ever - increasing cases of pathological sun reactions is one of the most important tasks of the dermatologist. Problems of pigmentation, physical and chemical sun protection, strength of the sun protection 'factor' (effective only up to 60% of the erythema threshold dose!) must be discussed with the patient exhibiting sun-induced skin lesions or just asking for advice.

L18 ANSWER 6 OF 6 DRUGU COPYRIGHT 2008 THE THOMSON CORP on STN

ACCESSION NUMBER: 1985-30177 DRUGU T M S Full-text

TITLE: Titanium Pigmentation: An Electron Probe

Microanalysis Study.

AUTHOR: Dupre A; Touron P; Daste J; Lassere J; Bonafe J L; Viraben R

LOCATION: Toulouse, France

SOURCE: Arch.Dermatol. (121, No. 5, 656-58, 1985) 6 Fig. 8 Ref.

CODEN: ARDEAC ISSN: 0003-987X

AVAIL. OF DOC.: Department de Dermatologie, Hopital de La Grave, Place Lange,

31052 Toulouse Cedex, France.

LANGUAGE: English

DOCUMENT TYPE: Journal FIELD AVAIL.: AB; LA; CT

FILE SEGMENT: Literature

N 1985-30177 DRUGU T M S Full-text

AB Following the use of a <u>titanium dioxide</u>-containing cream (Parkipan); other ingredients amyleine hydrate, trypan blue, white petrolatum, lanolin), for the treatment of herpetic erosive balanoposthitis, a patient developed a penile metallic pigmentation. Electron probe microanalysis showed that the pigmentation was probably due to deposition of titanium in the dermis facilitated by the presence of herpetic lesions.

ABEX A 22-yr-old man presented with asymptomatic, persistent, pearly, yellow pin-head sized shiny papules on the glans, sulcus and prepuce which had been present for 6 mth. The papules developed within 3 wk of starting treatment of herpetic lesions with Parkipan. Light microscopy showed normal skin morphology, except for the presence of brown granules in the upper part of the dermis. The granules were not in the epidermis or glandular structures. Polarized light microscopy failed to visualize the granules but dark field microscopy revealed refractive granules throughout the dermis. Under EM, the granules (500-600 nm diameter) were free in the dermis or located in macropahges, either free or within lysosome-like bodies. Electron probe microanalysis using energy dispersive analysis of X-ray (EDAX) films showed a major titanium peak for the pigment particles but not for surrounding tissues.

#### SEARCH HISTORY

### => d his ful

(FILE 'HOME' ENTERED AT 16:51:05 ON 22 JAN 2008)

FILE 'REGISTRY' ENTERED AT 16:59:46 ON 22 JAN 2008

4 SEA ABB=ON (SILVER OXIDE OR TITANIUM DIOXIDE OR ZINC OXIDE)/CN

L2 1 SEA ABB=ON (PIGMENTS OR INKS OR LACQUERS OR PLASTICS)/CN

FILE 'HCAPLUS' ENTERED AT 17:01:25 ON 22 JAN 2008

L3 288381 SEA ABB=ON L1 OR SILVER OXIDE OR TITANIUM DIOXIDE OR ZINC

L4 44662 SEA ABB=ON L3 AND (?PIGMENT? OR ?INK? OR ?LACQUER? OR

?PLASTIC?)
L5 17 SEA ABB=ON L4 AND ?HERPES?

E BUCHHOLZ HERWIG/AU

L6 102 SEA ABB=ON ("BUCHHOLZ HERWIG"/AU OR "BUCHHOLZ HERWIG A"/AU OR "BUCHHOLZ HERWING"/AU)

E BICARD BENHAMOU VALERIE/AU

L7 5 SEA ABB=ON "BICARD BENHAMOU VALERIE"/AU

E BRUNNER MARKUS/AU

23 SEA ABB=ON ("BRUNNER MARKUS"/AU OR "BRUNNER MARKUS DIPL
ING"/AU)

E MEDUSKI JERZY/AU

13 SEA ABB=ON ("MEDUSKI J"/AU OR "MEDUSKI J W"/AU OR "MEDUSKI JERZY"/AU OR "MEDUSKI JERZY D"/AU OR "MEDUSKI JERZY W"/AU)

L10 0 SEA ABB=ON L6 AND L7 AND L8 AND L9

L11 147 SEA ABB=ON L6 OR L7 OR L8 OR L9 L12 1 SEA ABB=ON L11 AND ?HERPES?

L13 47 SEA ABB=ON L11 AND ! HERFES!

FILE 'MEDLINE, BIOSIS, RAPRA, KOSMET, EMBASE, DRUGU' ENTERED AT 17:08:22
ON 22 JAN 2008

L16 24737 SEA ABB=ON L1 OR SILVER OXIDE OR TITANIUM DIOXIDE OR ZINC OXIDE

L17 5274 SEA ABB=ON L16 AND (?PIGMENT? OR INK? OR ?LACQUER? OR ?PLASTIC?)

L18 6 SEA ABB=ON L17 AND ?HERPES?

### FILE HOME

### FILE REGISTRY

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.  $\,$ 

STRUCTURE FILE UPDATES: 21 JAN 2008 HIGHEST RN 1000370-19-3 DICTIONARY FILE UPDATES: 21 JAN 2008 HIGHEST RN 1000370-19-3

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FILE COVERS 1907 - 22 Jan 2008 VOL 148 ISS 4 FILE LAST UPDATED: 21 Jan 2008 (20080121/ED)

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FILE LAST UPDATED: 19 Jan 2008 (20080119/UP). FILE COVERS 1949 TO DATE.

MEDLINE has been updated with the National Library of Medicine's revised 2008 MeSH terms. See HELP RLOAD for details.

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FILE BIOSIS

FILE COVERS 1926 TO DATE.

CAS REGISTRY NUMBERS AND CHEMICAL NAMES (CNs) PRESENT

FROM JANUARY 1926 TO DATE.

RECORDS LAST ADDED: 16 January 2008 (20080116/ED)

BIOSIS has been augmented with 1.8 million archival records from 1926 through 1968. These records have been re-indexed to match current BIOSIS indexing.

FILE RAPRA

FILE LAST UPDATED: 9 JAN 2008 <20080109/UP>

FILE COVERS 1972 TO DATE

>>> Simultaneous left and right truncation is available in the
basic index (/BI), and in the controlled term (/CT),
geographical term (/GT), and non-polymer term (/NPT) fields. <<</pre>

>>> The RAPRA Classification Code is available as a PDF file

>>> and may be downloaded free-of-charge from:

>>> http://www.stn-international.de/stndatabases/details/rapra\_classcodes.

FILE KOSMET

FILE LAST UPDATED: 2 JAN 2008 <20080102/UP>

FILE COVERS 1968 TO DATE.

>>> SIMULTANEOUS LEFT AND RIGHT TRUNCATION IS AVAILABLE IN THE BASIC INDEX (/BI) FIELD <><

#### FILE EMBASE

FILE COVERS 1974 TO 21 Jan 2008 (20080121/ED)

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>>> DERWENT DRUG FILE (SUBSCRIBER) <<<

- >>> FILE COVERS 1983 TO DATE <<<
- >>> THESAURUS AVAILABLE IN /CT <<<